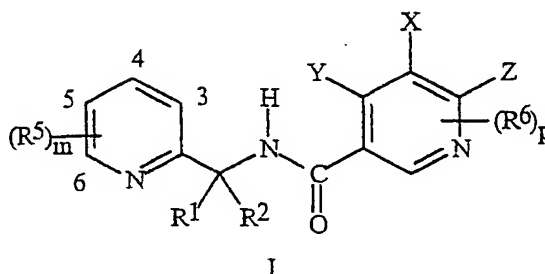


CLAIMS

What is claimed is:

1. A compound selected from Formula I, and *N*-oxides and agriculturally suitable salts thereof,



wherein

R^1 and R^2 are each independently H or C_1 - C_6 alkyl;

X and either Y or Z are a linking chain 3 or 4 atoms in length attached to contiguous carbon atoms and are taken together with said carbon atoms to form a fused phenyl ring, a fused 5- or 6-membered nonaromatic carbocyclic or heterocyclic ring optionally including one or two ring members selected from the group consisting of $C(=O)$, SO and $S(O)_2$, or a fused 5- or 6-membered heteroaromatic ring, each fused ring optionally substituted with one to three substituents independently selected from R^7 ;

each R^5 is independently C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, C_1 - C_6 haloalkyl, C_2 - C_6 haloalkenyl, C_2 - C_6 haloalkynyl, C_3 - C_6 halocycloalkyl, halogen, CN, CO_2H , $CONH_2$, NO_2 , hydroxy, C_1 - C_4 alkoxy, C_1 - C_4 haloalkoxy, C_1 - C_4 alkylthio, C_1 - C_4 alkylsulfinyl, C_1 - C_4 alkylsulfonyl, C_1 - C_4 haloalkylthio, C_1 - C_4 haloalkylsulfinyl, C_1 - C_4 haloalkylsulfonyl, C_1 - C_4 alkylamino, C_2 - C_8 dialkylamino, C_3 - C_6 cycloalkylamino, C_2 - C_6 alkylcarbonyl, C_2 - C_6 alkoxy carbonyl, C_2 - C_6 alkylaminocarbonyl, C_3 - C_8 dialkylaminocarbonyl or C_3 - C_6 trialkylsilyl;

each R^6 is independently C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, C_1 - C_6 haloalkyl, C_2 - C_6 haloalkenyl, C_2 - C_6 haloalkynyl, C_3 - C_6 halocycloalkyl, halogen, CN, CO_2H , $CONH_2$, NO_2 , hydroxy, C_1 - C_4 alkoxy, C_1 - C_4 haloalkoxy, C_1 - C_4 alkylthio, C_1 - C_4 alkylsulfinyl, C_1 - C_4 alkylsulfonyl, C_1 - C_4 haloalkylthio, C_1 - C_4 haloalkylsulfinyl, C_1 - C_4 haloalkylsulfonyl, C_1 - C_4 alkylamino, C_2 - C_8 dialkylamino, C_3 - C_6 cycloalkylamino, C_2 - C_6 alkylcarbonyl, C_2 - C_6 alkoxy carbonyl, C_2 - C_6 alkylaminocarbonyl, C_3 - C_8 dialkylaminocarbonyl or C_3 - C_6 trialkylsilyl;

each R⁷ is independently C₁-C₄ alkyl, C₂-C₄ alkenyl, C₂-C₄ alkynyl, C₃-C₆ cycloalkyl, C₁-C₄ haloalkyl, C₂-C₄ haloalkenyl, C₂-C₄ haloalkynyl, C₃-C₆ halocycloalkyl, halogen, CN, NO₂, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, C₃-C₆ cycloalkylamino, C₃-C₆ (alkyl)cycloalkylamino, C₂-C₄ alkylcarbonyl, C₂-C₆ alkoxycarbonyl, C₂-C₆ alkylaminocarbonyl, C₃-C₈ dialkylaminocarbonyl or C₃-C₆ trialkylsilyl;

m is 1, 2, 3 or 4; and

p is 0, 1, or 2.

2. The compound of Claim 1 wherein X and either Y or Z and the carbon atoms to which they are attached form a fused phenyl ring, a fused 5- or 6-membered nonaromatic carbocyclic ring or a fused 5- or 6-membered nonaromatic heterocyclic ring, each fused ring optionally substituted with one to three substituents independently selected from R⁷.

3. The compound of Claim 2 wherein one R⁵ is in the 3-position and a second R⁵ is in the 5-position and said two R⁵ groups are independently selected from the group consisting of C₁-C₆ alkyl, C₁-C₆ haloalkyl, halogen, CN, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, C₁-C₄ haloalkylthio, C₁-C₄ haloalkylsulfinyl and C₁-C₄ haloalkylsulfonyl.

4. The compound of Claim 3 wherein the R⁵ in the 3-position is selected from halogen and the R⁵ in the 5-position is selected from the group consisting of halogen, C₁-C₆ haloalkyl, C₁-C₄ haloalkoxy, C₁-C₄ haloalkylthio, C₁-C₄ haloalkylsulfinyl and C₁-C₄ haloalkylsulfonyl.

5. The compound of Claim 4 wherein the R⁵ in the 3-position is selected from halogen and the R⁵ in the 5-position is selected from the group consisting of halogen, C₁-C₆ haloalkoxy and C₁-C₆ haloalkyl.

6. The compound of Claim 5 wherein the R⁵ in the 3-position is chloro and the R⁵ in the 5-position is trifluoromethyl.

7. The compound of Claim 5 wherein the R⁵ in the 3-position is chloro and the R⁵ in the 5-position is selected from halogen or C₁-C₆ haloalkoxy.

8. The compound of Claim 1 wherein R¹ is H and R² is CH₃.

9. The compound of Claim 1 wherein each R⁶ is independently selected from the group consisting of halogen, C₁-C₆ alkyl and C₁-C₆ haloalkyl.

10. The compound of Claim 1 which is selected from the group:

2,4-dichloro-*N*-[[3-chloro-5-(trifluoromethyl)-2-pyridinyl]methyl]-5, 6, 7, 8-tetrahydro-3-quinolinecarboxamide;

N-[1-(5-bromo-3-chloro-2-pyridinyl)ethyl]-3-chloro-4-isoquinolinecarboxamide; and 3-bromo-*N*-[1-(5-bromo-3-chloro-2-pyridinyl)ethyl]-4-isoquinolinecarboxamide; and *N*-[1-(5-bromo-3-chloro-2-pyridinyl)ethyl]-3-fluoro-4-isoquinolinecarboxamide.

11. A fungicidal composition comprising a fungicidally effective amount of a compound of Claim 1 and at least one additional component selected from the group consisting of agriculturally suitable surfactants, solid diluents and liquid diluents.

5 12. A fungicidal composition comprising a fungicidally effective combination mixture of at least one compound of Claim 1 and at least one other fungicide.

13. The composition of claim 12 comprising (a) at least one compound of Formula I; and

(b) at least one compound selected from the group consisting of

10 (b1) alkylenebis(dithiocarbamate) fungicides;

(b2) compounds acting at the bc_1 complex of the fungal mitochondrial respiratory electron transfer site;

(b3) cymoxanil;

(b4) compounds acting at the demethylase enzyme of the sterol biosynthesis pathway;

15 (b5) morpholine and piperidine compounds that act on the sterol biosynthesis pathway;

(b6) phenylamide fungicides;

(b7) pyrimidinone fungicides;

(b8) phthalimides; and

(b9) fosetyl-aluminum.

20 14. The composition of claim 13 wherein the weight ratio of component (b) to component (a) is from 9:1 to 4.5:1.

15. A method for controlling plant diseases caused by fungal plant pathogens comprising applying to the plant or portion thereof, or to the plant seed or seedling, a fungicidally effective amount of a compound of Claim 1 or a composition thereof.

25 16. A method for controlling plant diseases caused by fungal plant pathogens comprising applying to the plant or portion thereof, or to the plant seed or seedling, a fungicidally effective amount of a composition of Claim 11.

17. A method of making the compound of Claim 10 consisting essentially of the procedure of Example 1.